RECEIVED **CENTRAL FAX CENTER**

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IN THE CLAIMS

1. (original) Uracils A uracil having general formula (I):

$$X_1$$
 X_2
 X_3
 X_4

(I)

wherein:

3 X_1 represents a hydrogen atom or a halogen atom;

- 4 X2 represents a halogen atom;
- X₄ represents a C₁-C₃ haloalkyl group;
- R represents a hydrogen atom, a C_1 - C_3 alkyl group or a C_1 - C_3 haloalkyl group;
- G represents an oxygen atom or a sulphur atom;
- X_3 represents a $Q(CR_1R_2)_nZ$ group, a Q_2 group, a Y(OC)- CR_6 = CR_5 - CR_3R_4Z group;
- Z represents an oxygen atom or a sulphur atom;
- R_1 , R_2 , R_3 and R_4 , the same or different, represent a hydrogen atom or, a C_1 - C_4 alkyl group; or a C_4 - C_4 haloalkyl group;
- R₅ represents an OR₇ group;
- R₆ represents a hydrogen atom or a C₁-C₄ alkyl group;
- R_7 represents a C_1 - C_4 alkyl group or a C_1 - C_4 haloalkyl group;
- Y represents a C₄-C₆ alkoxy or haloalkoxy group; an-OR₆

 group, a SR₉ group, a NR₁₀R₁₁ group;
- R₈ and R₉ represent a hydrogen atom, a C₁-C₆ linear or branched branched alkyl group, a C₁-C₆ linear or branched haloalkyl group, a C₂-C₆ eyeloalkyl group, a C₄-C₉ eyeloalkylalkyl group, a C₃-C₆ eyanoalkyl group, a C₃-C₆ alkonyalkyl group, an oxethonyl group, a tetrahydrofuranyl group; a phenyl group, a C₄-C₁₂ phenylalkyl group, a pyridyl group, said groups, in

turn, possibly substituted with one or more halogen atoms selected from chlorine, fluorine, bromine or iodine, or substituted with one or more groups selected from C₁-C₄-alkyl, or C₁-C₄-haloalkyl, C₁-C₄-alkoxy or C₄-C₄-haloalkoxy;

Rio and Rii, the same or different, represent a hydrogen atom, or a Ci Ci alkyl group, a Ci Ci haloalkyl group, a Ci Ci haloalkyl group, a Ci Ci exclealkyl group, a Ci Ci arylalkyl group, or an aryl group, said groups, in turn, possibly substituted with one or more halogen atoms selected from chlorine, fluorine, bromine or iodine, or substituted with one or more groups selected from a Ci Ci alkyl, or Ci Ci haloalkyl, Ci Ci alkoxy or Ci Ci haloalkyl, Ci Ci alkylene chain possibly substituted with Ci Ci alkyl groups and possibly interrupted by oxygen atoms or by a NRic group, wherein:

R₁₂ represents a hydrogen atom, a C₁-C₆ alkyl group or C₂-C₆ haloalkyl group, a C₂-C₆ alkenyl group or a C₃-C₆ haloalkenyl group, a C₂-C₆ alkynyl group or C₃-C₆ haloalkynyl group, a C₂-C₆ alkoxyalkyl group or a C₂-C₆ haloalkynyl group, a C₂-C₆ alkylearbonyl group or C₃-C₆ haloalkylearbonyl group:

- n represents 1, 2 or 3;
- Q represents a heterocyclic group selected from pyrrol—

2-yl, pyrrol 3-yl, imidazol-2-yl, imidazol-4-yl, imidazol-5-yl, pyrazol-3-yl, pyrazol-1-yl, pyrazol-5-yl, - 1,2,4-triazol-3-yl, - 1,2,4-triazol-5-yl, - 1,2,4- triazel-3-enyl, 1,2,3-triazelyl, tetrazelyl, exazelyl, isoxazol-5-yl, thiazol-2-yl, thiazol-5-yl, isothiazolyl, 1,3,4-oxadiazolyl, 1,3,4-thiadiazolyl, 1,2,4-thiadiazolyl, 1,2,4-oxadiazolyl, 1,2,4-oxadiazol-5-cn-3-yl, benzoxazol-2-yl, benzothiazol-2-yl, pyrazinyl, pyridazinyl, 1,2,4-triazinyl, 1,3,4thiadiazol-2-on-5-yl, 1,4,2-dioxazol-5-on-3-yl, 1,4,2oxathiazol-5-on-3-yl, 1,3,4-oxadiazin-5-on-2-yl, 1,4,2- $\frac{dio \times azin - 3 - yl}{dio \times azin - 3 - yl}$, $\frac{1, 2, 4 - o \times adiazin - 5 - on - 3 - yl}{dio \times azin - 3 - yl}$, $\frac{4, 5, 6, 7 - o \times adiazin - 5 - on - 3 - yl}{dio \times azin - 3 - yl}$ tetrahydro-1,3-benzothiazol-2-yl, 5,6-dihydro-4Heyelopenta[d][1,3]thiazele, said groups, in turn, possibly being optionally substituted with a halogen atom[[s]] selected from chlorine, fluorine, bromine or iodine, or substituted with a group[[s]] selected from C_1-C_6 alkyl or C_1-C_6 haloalkyl, C_2-C_6 alkenyl or C_2-C_6 haloalkenyl, 62-64 alkenyloxy or C2-C5 haloalkenyloxy, C_2-C_6 alkynyl or C_2-C_6 haloalkynyl, G_2-G_6 alkynyloxy or Ca-C6 haloalkynyloxy, C4-C6 alkoxy or C4-C6 haloalkoxy, C2-C6 alkoxyalkyl or C2-C6 haloalkoxyalkyl, C2-C6 alkoxyalkoxy, —G2-GE haloalkoxyalkoxy, haloalkoxyhaloalkoxy, C3-C3 alkoxyalkoxyalkyl, C3-C3

alkonyalkonyalkony, C1-C6 alkylthio or C1-C6 haloalkylthio, Ca-C6-alkylthioalkyl, C1-C6 alkylsulfinic or C1-C6 haloalkylsulfinic, C1-C6 alkylsulfonic or C1-C6 haloalkyloulfonic, C2-C6 alkoxycarbonyl or C2-C6 haloalkoxycarbonyl, C3-C2 alkenyloxycarbonyl or C3-C4 alkynyloxycarbonyl, Ca-Ca-alkoxycarbonylalkyl-or-Ca-Ca-Ca haloalkoxycarbonylalkyl, C4-C9 alkonylonycarbonylalkyl or G₄ G₂ alkynyloxycarbonylalkyl, C₂-C₃ alkoxycarbonylalkoxy, C4-Ca alkonyloxycarbonylalkoxy or G4-G2 alkynyloxycarbonylalkoxy, G2-G8 aminocarbonylalkoxy possibly substituted with G₁-G₄ alkyl groups or with a Co-Co alkylene group; GN, GHO, NO2, NH2, OH, C1-C2 cyanoalkyl, C1-C2 cyanoalkyloxy, C2-C6 formylalkyl, C2-C5 alkylcarbonyl, haloalkylcarbonyl, C2-C2 alkylcarbonylalkyl, C2-C6 alkonyimino, C2-C6 haloalkonyimino, C2-C6 alkoxyiminoalkyl, С₃-С₆ haloalkoxyiminoalkyl, С₃-С₆ alkoxyiminohaloalkyl, aminocarbonyl, C₂-C₆ aminocarbonylalkyl, aminoculfonyl or C2 C6 aminosulfonylalkyl, those last four groups possibly substituted with one or two Ci-C4 alkyl groups or with a G2-C5 alkylene group; C1-C6 alkyloulfonylamino, C2-C7 alkylcarbonylamino or C2-C2 alkoxycarbonylamino, these last three groups possibly substituted with Ca C4 alkyl

groups; C₆-C₁₀-aryl, C₅-C₁₂ arylalkyl, C₆-C₁₆-arylalkoxy, C₇-C₁₂ aryloxyalkyl, G₈-C₁₂ arylalkyloxyalkyl soid groups in turn possibly substituted with halogen atoms, C₁-C₄ alkyl groups, C₁-C₃_haloalkyl groups, C₁-C₄_alkoxy groups, C₁-C₃_haloalkoxy groups, CN; C₂-C₄_cycloalkyl, C₆-C₁₂_cycloalkylalkoxy, tetrahydropyran-2-yl said groups in turn possibly substituted with halogen atoms, C₄-C₄_alkyl groups, C₁-C₄_alkoxy groups;

Q_represents a heterocyclic group selected from

1,3,4-thiadiazol-2-yl, 1,3,4-thiadiazol-5-yl, 1,2,4thiadiazol-5-yl, tetrazol-5-yl, 1,3,4-exadiazol-2-yl,

1,3,4-exadiazol-5-yl, 1,2,4-exadiazol-5-yl, exazol-2yl, exazol-4-yl, exazol-5-yl, isexazol-3-yl, isexazol5-yl, thiazol-2-yl, thiazol-4-yl, thiazol-5-yl, said
groups, in-turn, possibly substituted with halogen
atoms selected from chlorine, fluorine, bromine er
iedine, or substituted with groups-selected from C₂-C₆
alkyl-er-C₁-C₆ haloalkyl, C₂-C₆ alkenyl-er-C₂-C₆
haloalkenyl, C₂-C₆ alkenylexy-er-C₂-C₆ haloalkenylexy,

C₂-C₆ alkynyl-er-C₂-C₆ haloalkynyl, C₂-C₆ alkynylexy-er
C₂-C₆ alkexyalkyl-er-C₂-C₆ haloalkexyalkyl, C₄-C₆
alkylthio-er-C₁-C₆ haloalkylthie, C₄-C₆ alkyleulfinic er

G1-G6 haloalkylsulfinie, G1-G6 alkylsulfonie or G1-G6 haloalkylsulfonic, C2-C6-alkoxycarbonyl or C2-C6 haloalkoxycarbonyl, Ca-Ca alkoxycarbonylalkyl or Ca-Ca haloalkoxycarbonylalkyl, C₂-C₆-alkoxycarbonylalkoxy, C₂-Caminocarbonylalkoxy possibly substituted with CarC4 alkyl-groups or with a Ga-Ca alkylene; - GN, CHO, NO2, NH2, C1-C2-cyanoalkyl, C1-C2-cyanoalkyloxy, C2-C4 alkylcarbonyl, -C2-C6-haloalkylcarbonyl, -C2-C5 alkoxyiminoalkyl, -C₂ -C₆ -haloalkoxyiminoalkyl, aminocarbonyl, C2-C6 aminocarbonylalkyl, aminoculfonyl o C2-C6 aminosulfonylalkyl, these last four groups possibly substituted with one or two C1 C4 alkyl groups or with a C2 C5 alkylono; C1 C5 alkyloulfonylamino, C2 C2 alkylearbonylamino or C2 C2 alkonycarbonylamino, these last three groups possibly substituted with C1-C4-alkyl groups; C4-C10 aryl, C6-C12 arylalkyl, C6-C10 arylalkoxy, C2-C12 aryloxyolkyl, C8-C12 arylalkyloxyalkyl said groups in turn possibly substituted with halogen atoms, C1-C4 alkyl-groups, C₁-C₂-haloalkyl groups, C₁-C₄-alkoxy groups, C1-C2 haloalkoxy groups, CN: C3 C2 eyeloalkyl, C6-C12 cycloalkylalkyl, C6-C16 cycloalkylalkoxy, tetrahydropyran-2-yl-said groups in turn possibly substituted with halogen atoms, C+-C+-alkyl groups, C+-C4_alkoxy-groups;

Q2 represents a heterocyclic group selected from 1Htetrazol-5-yl or 2H-tetrazol-5-yl, thiazol-2-yl, thiazol 4 yl, thiazol 5 yl, isothiazol 3-yl, isothiazol-4-yl, isothiazol-5-yl, 1,2,3-triazolyl, benzoxazol-2-yl, benzothiazol-2-yl, pyrimidin-2-yl, 1,2,4-triazinyl, 1,3,5-triazinyl, 1,3,4-thiadiazol-2on-5-yl, 1,4,2-dioxagol-5-on-3-yl, 1,4,2-oxathiazol-5on-3-yl, 1,3,4-oxadiazin-5-on-2-yl, 1,4,2-dioxazin-3v1, 1,2,4-oxodiazin-5-on-3-y1, 4,5,6,7-tetrahydro-1,3benzothiazol-2-yl, 5,6-dihydro-4Heyelopenta[d][1,3]thiazole, said groups in turn possibly being optionally substituted with halogen atoms scleeted from chlorine, fluorine, bromine or iodine, or substituted with a group[[s]] selected from: C_1-C_6 alkyl; [[or]] C_1-C_6 haloalkyl[[,]]; C_2-C_6 alkenyl; [[or]] C_2-C_6 haloalkenyl[[,]]; C_2-C_6 alkenyloxy or C_2-C_6 haloalkenyloxy, C2-C6 alkynyl; [[or]] C2-C6 haloalkynyl, G2-G6 alkynyloxy or C2-C6 haloalkynyloxy, G1-C6 alkoxy or C₁-C₆ haloalkoxy, C₂-C₆ alkoxyalkyl; [[or]] C₂-C₆ haloalkoxyalkyl[[,]]; Ca-Ca-alkoxyalkoxy, Ca-Ca haloalkoxyalkoxy, C2-C6 haloalkoxyhaloalkoxy, C3-C8 alkoxyalkoxyalkyl, C3-C8-alkoxyalkoxyalkoxy, C4-C6 alkylthio or C1-C6 haloalkylthio, C2-C6 alkylthioalkyl, C1-C6_alkylsulfinic or C1-C6_haloalkylsulfinic, C1-C6

alkyloulfonic or C1-C6 haloalkyloulfonic, C2-C6 alkoxycarbonyl or C2-C5 haloalkoxycarbonyl, C3-C1 alkenyloxyearbonyl or Ca-Calkynyloxycarbonyl, Ca-Ca alkowycarbonylalkyl or C3-C2 haloalkowycarbonylalkyl, C4-C2-alkenylonyearbonylalkyl or -C4-C2 alkynyloxycarbonylalkyl, G2-C8 alkoxycarbonylalkoxy, alkenyloxycarbonylalkoxy-C4-C4-c4-c4 alkynyloxycarbonylalkoxy C4-C4-C4-C4 aminocarbonylalkony possibly substituted with C1-C4 alkyl or with a C2-C5-alkylone; CN, CHO, NO2,-NH2,-OH, C1-C2 cyanoalkyl, C1-C2 cyanoalkyloxy, C2-C6 formylalkyl, C2-C4 alkylcarbonyl, C2 C6 haloalkylcarbonyl, C2-C2 alkylcarbonylalkyl, C2-C6 alkoxyimino, C2-C6 haloalkoxyimino, -C₂-C₅ alkoxyiminoalkyl, C₂-C₅ haloalkonyiminoalkyl, alkonyiminohaloalkyl Cz Co, aminocarbonyl, C2 C6 aminocarbonylalkyl, aminosulfonyl or C2 C6 aminoculfonylalkyl, these last four groups possibly substituted with one or two C1 C4 alkyl groups or with a C2-C5 alkylene; C.-C6 alkyloulfonylamino, C2-C2 alkylcarbonylamino o Ca-Ca-alkoxycarbonylamino, these last three groups possibly substituted with -G₁-G₄ alkyl groups; C₆-C₁₀ aryl, C₆-C₁₂ arylalkyl, C₆-C₁₀-arylalkoxy, C_7-C_{12} aryloxyalkyl, C_8-C_{12} arylalkyloxyalkyl said groups in turn possibly being optionally substituted with

halogen atoms, C_1 - C_4 alkyl groups, C_1 - C_3 haloalkyl groups, C_1 - C_4 alkoxy groups, C_1 - C_3 haloalkoxy groups, C_1 : C_3 - C_7 cycloalkyl, C_6 - C_{12} cycloalkylalkyl, C_6 - C_{10} cycloalkylalkoxy, tetrahydropyran-2-yl said groups in turn possibly being optionally substituted with halogen atoms, C_1 - C_4 alkyl groups, C_1 - C_4 alkoxy groups.

- 2.(original): The uracils A uracil according to claim 1, characterized in that they are it is selected from:
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-methoxybut-2-enoate;
- methyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-methoxybut-2-enoate;
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenylthio}-3-methoxybut-2-enoate;
- ethyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-ethoxybut-2-enoate;
- methyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenylthio}-3-methoxybut-2-enoate;
- ethyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-

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methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-ethoxybut-2-enoate;
- isopropyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-
tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-
1-y1)phenoxy)-3-methoxybut-2-enoate;
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-
2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-
methoxybut-2-enoate;
- methyl (2E)-4-\{2,4-\text{dichloro}-5-\{1,2,3,6-\text{tetrahydro}-2,6-
dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-
methoxybut-2-enoate;
- ethyl (2E)-4-\{2-chloro-4-fluoro-5-\{1,2,3,6-tetrahydro-
2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenoxy)-3-
ethoxybut-2-enoate;
 <del>-cthyl (25)-4-{2,4-dichloro-5-[1,2,3,6-tetrahydro-2,6-</del>
dioxo-1-(trifluoromethyl)pyrimidin l yl]phonoxy)
ethoxybut-2-enoate;
-2,2,2 trifluoroethyl-(2E)-4-(2 chloro-4-fluoro-5-
{1,2,3,6 tetrahydro-2,6-dioxo-1 (trifluoromethyl)pyrimidin-
1 yl]phenoxy) - 3 -methoxybut - 2 - encate >
- (2E)-4-(2-chloro-4-fluoro-5-(1,2,3,6-tetrahydro-2,6-
dioxo 1 (trifluoromethyl)pyrimidin l yl]phenoxy}-3-methoxy-
N, N-dimethylbut-2-enamide;
 S othyl (25) 1 (2 chlore 1 fluore 5 [1,2,3,6 tetrahydre
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2,6 diexo-1-(trifluoromethyl)pyrimidin-1-yl]phonoxy}-3-
methoxybut-2-enethioate;
- isopropyl (2E)-4-\{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-]
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxybut-2-enoate;
- 2,2,2-trifluoroethyl (2E)-4-{2-chloro-4-fluoro-5-
[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluorome-
thyl)pyrimidin-1-yl]phenoxy}-3-methoxybut-2-enoate;
- 2,2,2-trifluoroethyl (2E)-4-{2,4-dichloro-5-[1,2,3,6-
tetrahydro-3-methyl-2, 6-dioxo-4-(trifluoromethyl)
pyrimidin-1-yl]phenoxy}-3-methoxybut-2-enoate;
- S-ethyl (2E)-4-(2-chloro-1-fluoro-5-[1,2,3,6-tetrahydro
<del>3 methyl-2,6-dioxe-4-(trifluoromethyl) pyrimidin-1-</del>
yl]phonoxy]-3-methoxybut-2-enethioate;
 -S-ethyl (2E)-4-{2,4-dichloro-5-{1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxybut-2-enethioate>
- (2E)-4-(2-chloro-4-fluoro-5-(1,2,3,6-tetrahydro-3-methyl-
2,6 diono 1 (trifluoromethyl) pyrimidin 1 yl]phonoxy) 3
methoxy-N,N-dimethylbut-2-enamide;
-- (25)-4-(2,4-dichloro-5-(1,2,3,6-tetrahydro-3-methyl-2,6-
dioxo-1-(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-methoxy-
N, N dimethylbut 2 enamide;
  (25)-4-(2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-methyl-
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2,6-dioxo-1-(trifluoromethyl)pyrimidin-1-yl}phenylthio}-3-
methoxy-N, N-dimethylbut-2-enamide;
- (2E)-1-(2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-
dioxo 1 (trifluoromethyl)pyrimidin-1-yllphenylthio -3-
methoxy-N, N-dimethylbut-2-enamide;
- 3-[4-chlore-2-fluore-5-(tetrazel-5-ylmethexy)phenyl]-6-
(trifluoromethyl) 2, 4(1H, 3H) pyrimidinedione,
-3-{4-chloro-2-fluoro-5-{(2-methyl-2H-tetrazol-5-
v1)methoxy]pheny1) 6 (trifluoromethy1) 2, 4(1H, 3H)
pyrimidinedione;
- 3-[4-chlore-2-fluore-5-(tetrazel-5-ylmethoxy)phenyl] 1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedioner
- 3-[2,4-dichloro-5 (tetrazol 5 ylmethoxy)phonyl] l methyl-
6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
--3-(4-chloro-2-fluoro-5-[(2-methyl-2H-tetrazol-5-
yl)methoxylphenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
 3-{4-chlore-2-fluore-5-{(2-cthyl-2W-tetrazel-5-
v1) methoxy] phenyl] -1-methyl 6 (trifluoromethyl) 2, 4(1H, 3H) -
pyrimidinedione+
- 3-(2,4-dichlore-5-[(2-methyl-2H-tetrazol-5-
yl) methoxy] phenyl} -1-methyl -6-(trifluoromethyl) -2, 4(1H, 3H) -
pyrimidinedione;
- 3-(2,4-dichloro-5-((2-ethyl-2H-tetrazol-5-
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y1)methoxy]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
--3-{4-chloro-2-fluoro-5-[(1-cthyl-1H-tetrazol-5-
y1)methoxy]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione.
 -3 {2,4-dichloro-5-[(1-cthyl-1H-tetrazol-5-
y1)methony]phenyl]-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-{5-[(5-tert-butyl-1,3,4-oxadiazol-2-yl)methoxy}-4-
chloro-2-fluorophenyl}-1-methyl-6-(trifluoromethyl)-
2,4(1H,3H)-pyrimidinedione;
- methyl [5 ({2 chloro 1 fluoro 5 [1,2,3,6 tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}methyl}-1H-tetrazol-1-yl]acetate;
- methyl [5 ({2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-
2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy)methyl)-lH-tetrazol-l-yl]acetate;
-methyl-[5-([2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
mothyl-2,6-diono-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}methyl)-2H-tetrazol-2-yl}acetate;
-- methyl--[5-((2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-
2,6-diexe-4-(trifluoromethyl)pyrimidin-1-
yl-phenoxy)methyl) -2H-tetrazol-2-yl]acetate;
- 3-[4-chloro-3-(tetrazol-5-yl)phenyl]-6-(trifluoromethyl)-
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2,4(1H,3H)-pyrimidinedione;
 - 3-[4-chloro-3-(2-methyl-2H-tetrazol-5-yl)phenyl]-6-
 (trifluoromethyl) -2,4(1H,3H)-pyrimidinedione;
 - 3-[4-chloro-3-(1-methyl-1H-tetrazol-5-yl)phenyl]-6-
 (trifluoromethyl) -2,4(1H,3H)-pyrimidinedione;
 - 3-[4-chloro-3-(tetrazol-5-yl)phenyl]-1-methyl-6-
 (trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
 - 3-[4-chloro-2-fluoro-5-(tetrazol-5-yl)phenyl]-6-
 (trifluoromethyl) -2, 4 (1H, 3H) -pyrimidinedione;
 - 3-[2,4-dichloro-5-(tetrazol-5-yl)phenyl]-6-
 (trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(tetrazol-5-yl)phenyl]-1-methyl-6-
 (trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
-3-[2,4-dichloro-5-(tetrazol-5-yl)phenyl]-1-methyl-6-
 (trifluoromethy1) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[4-chloro-3-(2-methyl-2H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(2-methyl-2H-tetrazol-5-
y1) pheny1]-6-(trifluoromethy1)-2,4(1H,3H)-pyrimidinedione;
-3-[2,4-dichloro-5-(2-methyl-2H-tetrazol-5-yl)phenyl]-6-
(trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(1-methyl-1H-tetrazol-5-
yl)phenyl]-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
-3-[2,4-dichloro-5-(1-methyl-1H-tetrazol-5-yl)phenyl]-6-
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(trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(2-methyl-2H-tetrazol-5-
yl) phenyl] -1-methyl-6-(trifluoromethyl) -2, 4(1H, 3H) -
pyrimidinedione;
-3-[2,4-dichloro-5-(2-methyl-2H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- 3-[4-chloro-3-(2-ethy1-2H-tetrazol-5-yl)phenyl]-1-methyl-
6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
-3-[4-chloro-3-(1-methyl-1H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
-3-[4-chloro-2-fluoro-5-(1-methyl-1H-tetrazol-5-
yl) phenyl] -1-methyl-6-(trifluoromethyl) -2, 4(1H, 3H)-
pyrimidinedione;
-3-[2,4-dichloro-5-(1-methyl-1H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
-3-[4-chloro-3-(1-ethyl-1H-tetrazol-5-yl)phenyl]-1-methyl-
6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
- methyl (5 {2-chlore-5-{1,2,3,6-tetrahydre-3-methyl-2,6-
dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenyl}-1H-
tetrazol-1-yl)acetate;
--methyl (5-{2-chloro-5-{1,2,3,6-tetrahydro-3-methyl-2,6-
dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenyl}-2H-
tetrazol-2-yl)acetate;
methyl (5-{2-chloro-4-fluoro-5-{1,2,3,6-tetrahydro-3-
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methyl-2,6-dioxo-4 (trifluoromethyl)pyrimidin-1-yl]phenyl}-
1H-tetrazol-1-yl)acetate;
-methyl (5-[2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-1-(trifluoromethyl)pyrimidin-1-yl]phonyl}-
2H-tetrazol-2-yl)acetate;
-- methyl (5-{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-
2,6-dioxo 4-(trifluoromethyl)pyrimidin-1-yl]phenyl}-1H-
tetrazel-1-yl)acetate;
- mothyl-(5-[2,4-dichloro-5-[1,2,3,6-tetrahydro-3-methyl-
2,6-dioxe-4-(trifluoromethyl)pyrimidin-1-yl]phenyl]-2H-
tetrazol-2-yl) acetate;
-3-[4-chloro-3-(4-methoxy-5-methyl-1,3-thiazol-2-
vl) phenyl 6 (trifluoromethyl) 2, 4 (1H, 3H) pyrimidinedioner
 -3-[2,4-dichlore-5-(4-methoxy-5-methyl-1,3-thiazol-2-
v1) phenyl-6-(trifluoromethyl)-2, 4(1H, 3H)-pyrimidinedione;
- 3-[4-chloro-2-fluoro-5-(4-methoxy-5-methyl-1,3-thiazol-2-
v1) phenyl-6-(trifluoromethyl)-2, 4(1H, 3H) - pyrimidinedione;
-3-[4-chloro-3-(4-methoxy-5-methyl-1,3-thiazol-2-
yl) phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
-3-[4 chloro-3 (4-cthoxy-5-methyl-1,3-thiazol-2-yl)phenyl-
1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
 3 [2,4-dichloro-5-(4 methoxy 5-methyl-1,3-thiazol-2-
yl) phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
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pyrimidinedione;
-3-[2,4-dichlore-5-(4-ethoxy-5-methyl-1,3-thiazel-2-
vl) phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
eyrimidinedione;
-3-[4-chloro-2-fluoro-5-(4-methoxy-5-methyl-1,3-thiazol-2-
v1) phonyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
-3-[4-chloro-2-fluoro-5-(4-cthoxy-5-mcthyl-1,3-thiazol-2-
yl) phonyl-1-methyl-6-(trifluoremethyl)-2,4(1H,3H)-
pyrimidinedione;
-3-{4-chlore-3-(4-benzylexy-5-methyl-1,3-thiazel-2-
yl)phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-[2,4-dichlore-5-(4-benzyloxy-5-methyl-1,3-thiazel-2-
yl) phenyl-1-mothyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
-3-[4-chlore-2-fluore-5-(4-benzyloxy-5-methyl-1,3-thiazel-
2-yl)phenyl-1-methyl-6-(trifluoromethyl) 2,4(1H,3H)-
pyrimidinediono;
-3-(2,4-dichlore-5-([5-(trifluoromethyl)-1,3,4-thiadiazol-
2-yl]oxy|phenyl)-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione+
- 3-(4-chloro-2-fluoro-5-({5-(trifluoromothyl)-1,3,4-
thiadiazol-2-yl]oxy)phenyl)-6-(trifluoromethyl)-2,4(1H,3H)-
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pyrimidinediene;
-3-(2,4-dichlore-5-({5-(trifluoromethyl)-1,3,4-exadiazel-
2-yl]ony)phenyl)-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
-3-(4-chlore-2-fluore-5-([5-(trifluoremethyl)-1;3,4-
oxadiazel-2-yl]oxy}phenyl)-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedioner
-3-(4-chloro-3-([5-(trifluoromethyl)-1,3,4-thiadiazel-2-
yljoxy)phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-(2,4-dichloro-5-([5-(trifluoromethyl)-1,3,4-thiadiazol-
2-yl]oxy]phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-(4-ehlore-2-fluore-5-([5-(trifluoremethyl)-1,3,4-
thiadiazol-2-yl]oxy}phenyl)-1-methyl-6-(trifluoromethyl)-
2,4(1H,3H)-pyrimidinedione;
- 3-{4-chlore-3-[(5-methyl-1,3,4-thiadiazel-2-
yl) oxylphenyl} - 1 - methyl - 6 - (trifluoromethyl) - 2, 4 (1H, 3H) -
pyrimidinedione;
-3-{2,4-dichloro-5-[(5-methyl-1,3,4-thiadiazol-2-
yl) oxylphenyl}-1-methyl-6 (trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
 -3-{4-chloro-2-fluoro-5-{-(5-methyl-1,3,4-thiadiazol-2-
yl) oxy]phenyl]-1-methyl-6 (trifluoromethyl)-2,4(1H,3H)-
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```
pyrimidinedione;
-3-(4-chloro-3-([5-(trifluoromethyl)-1,3,4-oxadiazol-2-
yl]oxy)phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
 -3-(2,4-dichlore-5-([5-(trifluoromethyl)-1,3,4-oxadiazol-
2-y1]oxy]phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedioner
-3-(4-chlore-2-fluore-5-(-5-(trifluoremethyl)-1,3,4-
oxadiazol-2-yl]oxy)phenyl)-1-methyl-6-(trifluoromethyl)-
2,4(1H,3H) -pyrimidinedione+
- 3-(4-chlore-3-[(5-methyl-1,3,4-exadiazel-2-
yl) oxylphonyl) -1-methyl-6 (trifluoromethyl) -2,4(1H,3H)-
pyrimidinedione;
-3-{2,4-dichloro-5-{\(\frac{5}{methyl-1,3,4-oxadiazol-2-\)}}
yl)oxy]phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-(4 ehloro-2 fluoro-5 (5-methyl-1,3,4-emadiazol-2-
yl)oxy]phenyl]-1-methyl-6 (trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-6-oxo-2-thioxo-4-(trifluoromethyl)pyrimidin-1-
y1]phenoxy}-3-methoxybut-2-enoate;
- methyl (2E)-4-\{2-chloro-4-fluoro-5-\{1,2,3,6-tetrahydro-3-
difluoromethyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
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```
yl]phenoxy}-3-methoxybut-2-enoate;
    3-[4-chlore-3-(4,5-dimethyl-1,3-thiazol-2-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione+
              (2E)-4-\{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-
     methyl
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxypent-2-enoate;
- methyl (2E)-4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxypent-2-enoate;
      ethy1 (2E)-4-\{2,4-dichloro-5-[1,2,3,6-tetrahydro-3-
methy1-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxybut-2-enoate;
- ethyl (2E) -4-{2-chloro-4-fluoro-5-[1,2,3,6-tetrahydro-3-
methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-
yl]phenoxy}-3-methoxybut-2-enoate;
             3-\{4-\text{chloro}-3-[2-(\text{methoxymethyl})-2H-\text{tetrazol}-5-
yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1.H, 3.H)-
pyrimidinedione;
              3-4-chloro-3-[1-(methoxymethyl)-1H-tetrazol-5-
yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
   3-\{4-\text{chloro}-3-[2-(\text{ethoxymethyl})-2H-\text{tetrazol}-5-yl]phenyl}-
1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
 -3-{4-chloro}-3-[1-(ethoxymethyl)-1H-tetrazol-5-yl]phenyl}-
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```
1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
   3-[3-(2-allyl-2H-tetrazol-5-yl)-4-chlorophenyl]-1-methyl-
6-(trifluoromethyl) - 2,4(1H,3H)-pyrimidinedione;
- 3-[3-(1-allyl-1H-tetrazol-5-y1)-4-chlorophenyl]-1-methyl-
6-(trifluoromethyl) - 2,4(1H,3H)-pyrimidinedione;
               3-{4-chlore-2-fluore-5-{(3-methylisexazel-5-
y1) methoxy]phenyl} 1-methyl-6-(trifluoromethyl)-2,4-(1H,3H)-
pyrimidinedione+
                     3-{2,4-dichlero-5-{-(3-methylisoxasol-5-
yl) methoxy | phenyl | -1-mothyl -6-(trifluoromethyl) -2, 4 (1H, 3H) -
pyrimidinedione:
       -3-[4-chloro-3-(4-isopropoxy-5-methyl-1,3-thiczol-2-
+1) phenyl-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinediene;
           -3-[4-chloro-3-(4-hydroxy-5-methyl-1,3-thiazel-2-
yl) phenyl-1 methyl-6-(trifluoromethyl)-2, 4(1H, 3H)-
pyrimidinedione;
        3-{4-chloro-2-fluoro-5-[(5-methyl-1,2,4-oxadiazol-3-
y1) methoxy]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
              3-{2,4-dichloro-5-[(5-methyl-1,2,4-oxadiazol-3-
y1) methoxy] pheny1}-1-methy1-6-(trifluoromethy1)-2, 4(1H, 3H)-
pyrimidinedione;
   3-{3-(1,3-benzethiazol-2-yl)-4-chlorophenyl}-1-methyl-6-
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```
(trifluoromethyl) - 2,4(1H,3H) - pyrimidinedione;
     -3-[3-<del>(1,3-benzoжazol-2-yl)-4-chloxophenyl]-1-methyl-6-</del>
(trifluoromethyl) - 2,4(1H,3H) pyrimidinediones
        3-{4-chloro-2-fluoro-5-[(3-methyl-1,2,4-oxadiazol-5-
yl)methoxylphenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
  -3-[4-chlore-3-(4-methyl-1,3-thiazel-2-yl)phenyl-1-methyl-
6-(trifluoromethyl)-2, 4(1H, 3H)-pyrimidinedione+
                   3-[4-chloro-2-fluoro-5-(1,2,4-oxadiazo].-3-
ylmethoxy) phenyl] -1-methyl-6-(trifluoromethyl) -2, 4(1H, 3H) -
pyrimidinedione;
     3-[3-(2-tert-butyl-2H-tetrazol-5-yl)-4-chlorophenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
   - 3-[5-(1,3-benzothiazel-2-yl)-4-chlore-2-fluorophenyl]-1-
methyl-6-(trifluoromethyl)- 2,4(1H,3H)-pyrimidinedione;
-3-(4-\text{chloro}-3-(2-[(2-\text{methoxyethoxy})\text{methyl}]-2H-\text{tetrazol}-5-
yl)phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
- 3-(4-chloro-3-{1-[(2-methoxyethoxy)methyl]-1H-tetrazol-5-
y1) pheny1) -1-methy1-6-(trifluoromethy1) -2, <math>4(1H, 3H) -
pyrimidinedione;
   3-(5-(1,3-benzoxazel-2-yl)-4-chlore-2-fluorophenyl}-1-
methyl 6 (trifluoromethyl) - 2,4(1H,3H) - pyrimidinedione;
        _ 3-[5-(1,3-benzothiazol-2-yl)-2,4-dichlorophenyl]-1-
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yl) methoxy] phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
      3-[4-chloro-3-(2-isopropyl-2H-tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
          3-[3-(2-benzyl-2H-tetrazol-5-yl)-4-chlorophenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
          3-[3-(1-benzyl-1H-tetrazol-5-yl)-4-chlorophenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
            -3-{4-chlore-2-fluore-5-{(1-methyl-1H-tetrazel-5-
yl) oxylphenyl}-1-methyl 6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
         ____3-(4-chlore-2-fluore-5-[(2-methyl-2H-tetrazel-5-
yl) oxylphenyl | 1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione/
    methyl (2E)-4-\{2-\text{chloro}-5-[1,2,3,6-\text{tetrahydro}-3-\text{methyl}-\text{methyl}-\text{methyl}-\text{methyl}-\text{methyl}
2,6-dioxo-4(trifluoromethyl)pyrimidin-1-yl]phenoxy}-3-
methoxybut-2-enoate;
            (2E)-4-\{2-\text{chloro}-5-[1,2,3,6-\text{tetrahydro}-3-\text{methyl}-
2,6-dioxo-4(trifluoromethyl)pyrimidin-1-y1]phenoxy}-3-
ethoxybut-2-enoate;
        3-[4-chloro-3-(1,2,4-oxadiazol-3-ylmethoxy)phenyl]-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
   3 (4-chlore-3 (3-methylisexazel-5-yl)methexylphenyl)-1-
mothyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
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```
-3-[4-ehloro-3-(4,5,6,7-tetxahydro-1,3-benzethiazol-2-
yl) phenyl]-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione:
  -3-[4-chloro-3-(5,6-dihydro-1,4,2-dioxazin-3-yl)phenyl}-1-
methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
       -3-[4-chlore-3-(4-methyl-5-exe-5,6-dihydre-4H-1,3,4-
oxadiazin 2-yl)phonyl}-1-methyl-6-(triflueromethyl)-
2,4(1H,3H)-pyrimidinedione;
_3-{4-chlore-3-(5,6-dihydro-1,4,2-dioxazin-3-ylmethoxy)-2-
fluorophenyl]-1-methyl-6-(trifluoromethyl)-2,4-(1H,3H)-
pyrimidinedione;
    -3-{4-chloro-2-fluoro-5-[-(4-methyl-5-exe-5,6-dihydre-4#-
1,3,4-oxadiazin-2-yl)methoxy]phonyl)-1-methyl-6-
(trifluoremethyl)-2,4(1H,3H)-pyrimidinedione;
       3-[4-chloro-3-(2-phenyl-2H-tetrazol-5-yl)phenyl]-1-
methyl-6 (trifluoremethyl)-2,4(1H,3H)-pyrimidinedioner
       3-[4-chlore-3-(1-phenyl-1H tetrazol-5-yl)phenyl]-1-
methyl-6-(trifluoremethyl)-2,4(1H,3H)-pyrimidinediene;
         3-{4-chloro-3-[1-(cyclopropylmethyl)-1H-tetrazol-5-
yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
         3-{4-chloro-3-[2-(cyclopropylmethyl)-2H-tetrazol-5-
yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
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```
<del>_ 3 (4 chloro 3 [1 (2 exopropyl) 1 H-tetrazel 5 yl]phenyl} -</del>
1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
  3-44-chlore-3-{2-(2-oxopropyl)-2H-tetrarel-5-yl]phonyl}-
1-methyl 6 (trifluoromethyl) -2, 4(1H, 3H) -pyrimidinedione;
 -3-[4-chlore-3-(4-cyclopropyl-1,3-thiazel-2-yl)phenyl]-1-
methyl -6-(trifluoromethyl)-2, 4(1H, 3H)-pyrimidinedioner
          3-{4-chloro-3-{4-(4-chlorophenyl)-1,3-thiazol-2-
yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
   cthyl 2-(2-chlore-5-[1,2,3,6-tetrahydre-3-methyl-2,6-
dioxo-1-(trifluoromothyl)pyrimidin 1-yl]phonyl)-1,3-
thiazole 4 carboxylater
- 3-[3-(2-butyl-2H-tetrazol-5-yl)-4-chlorophenyl]-1-methyl-
6-(trifluoromethyl)-2,4(1H,3H)-pyrimidinedione;
-3-[4-chloro-2-fluoro-5-(5,6-dihydro-1,4,2-dioxazin-3-
ylmethoxy) 2-fluorophenyl]-1-methyl-6-(trifluoromethyl)-
2,4(1H,3H)-pyrimidinedione;
- 3-(4-\text{chloro}-3-\{2-[(4-\text{chlorophenoxy})\text{methyl}]-2H-\text{tetrazol}-5-
y1}phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
   3-(4-chloro-3-(1-[(4-chlorophenoxy)methyl]-1H-tetrazol-5-
yl}phenyl)-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)-
pyrimidinedione;
   3-[3-(4-text-butyl-5-0x0-4,5-dihydro-1,3,4-thiadiazol-2-
```

yl)-4-chlorophenyl]-1-methyl-6-(trifluoromethyl)2,4(1H,3H)-pyrimidinedione;

- 3-{4-chloro-3-[2-(4-chlorobenzyl)-2H-tetrazol-5-yl]phenyl}-1-methyl-6-(trifluoromethyl)-2,4(1H,3H)pyrimidinedione;
- $-3-\{4-\text{chloro}-3-[1-(4-\text{chlorobenzyl})-1H-\text{tetrazol}-5-yl] \\ \text{phenyl}-1-\text{methyl}-6-(\text{trifluoromethyl})-2, 4(1H, 3H)-\\ \text{pyrimidinedione};$
- methyl 2-{2-chloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenyl}-1,3thiazole-4-carboxylate;
- methyl (2-{2-chloro-5-[1,2,3,6-tetrahydro-3-methyl-2,6-dioxo-4-(trifluoromethyl)pyrimidin-1-yl]phenyl}-1,3-thiazol-4-yl)acetate.
- (canceled)
- 4. (withdrawn): A process for the preparation of compounds having general formula (I) according to claim 1, characterized in that it includes a cyclo-condensation reaction of an isocyanate or isothiocyanate having general formula (II) with a 3-aminocrotonate having general formula (III) according to reaction scheme 1

 Scheme 1:

NCG
$$X_4$$
 X_2
 X_3
 X_4
 X_4
 X_4
 X_4
 X_5
 X_4
 X_4
 X_5
 X_5
 X_6
 X_4
 X_5
 X_7
 X_8
 X_9
 X_9

wherein

- X_1 , X_2 , X_3 , X_4 , R and G have the meanings previously defined;
- R_{13} represents a C_1 - C_4 alkyl or C_1 - C_4 haloalkyl group or a phenyl group possibly substituted with C_1 - C_4 alkyl groups.
- 5. (withdrawn): The process according to claim 4, characterized in that the reaction is carried out in the presence of an inert organic solvent and in the presence of an organic base or preferably inorganic base, at a temperature ranging from -20°C to the boiling point of the reaction mixture.
- 6.(withdrawn): The process according to claim 4, characterized in that the isocyanates or isothiocyanates having general formula (II) are prepared starting from a

substituted aniline having general formula (IV) by reaction with a compound having general formula (V), such as phosgene, diphosgene, triphosgene or thiophosgene, according to reaction scheme 2

Scheme 2:

wherein

- X_1 , X_2 , X_3 and G have the meanings defined above;
- L_3 and L_4 , the same or different, represent a chlorine atom or a CCl₃O- group.
- 7. (withdrawn): The process according to claim 6, characterized in that the reaction is carried out in the presence of an inert organic solvent, at a temperature ranging from 0°C to the boiling point of the mixture itself, possibly in the presence of a catalyst such as triethylamine, in an amount ranging from 0.001 and 100% by weight with respect to the aniline (IV), with a quantity of

reagent (V) varying from 1 to 3 moles per mole of aniline (IV).

8. (withdrawn): The process for the preparation of compounds having general formula (I) according to claim 1, wherein X_3 represents a $Q(CR_1R_2)_nZ$ - group, a Q_1Z - group, a Y(OC)- CR_6 = CR_5 - CR_3R_4Z - group, compounds (Ia), characterized in that it comprises the reaction of a uracil having general formula (VI) with a compound having general formula (VII) according to reaction scheme 3

$$X_1$$
 X_2
 X_3
 X_4
 X_4
 X_4
 X_4
 X_4
 X_5
 X_4
 X_5
 X_4
 X_5
 X_6
 X_7
 X_8
 X_8
 X_9
 X_9

Scheme 3:

wherein

- X_1 , X_2 , X_4 , G and Z have the meanings previously defined;
- R represents a C_1-C_3 alkyl group or a C_1-C_3 haloalkyl group;
- W represents a $Q(CR_1R_2)_n$ group, a Q_1 group, a Y(OC)- CR_6 = CR_5 - CR_3R_4 group, wherein R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , Y, Q and Q_1 have the meanings defined above;
- L_2 represents a halogen atom, a R_LSO_2O- group, wherein R_L represents a C_1-C_4 alkyl or C_1-C_4 haloalkyl group or a phenyl group possibly substituted by C_1-C_4 alkyl groups, or it represents a $R_{L1}SO_2-$ group wherein R_{L1} represents a C_1-C_4 alkyl or C_1-C_4 haloalkyl group.
- 9. (withdrawn): The process according to claim 8, characterized in that the reaction between the compounds having general formula (VI) and the compounds having general formula (VII) is carried out in the presence of one or more inert organic solvent(s) and in the presence of a base, preferably an inorganic base, at a temperature ranging from -10°C to the boiling point of the reaction mixture.
- 10. (withdrawn): The process for the preparation of the compounds having general formula (I) according to claim 1, wherein G = O and $R \neq H$, compounds (Ic), characterized in that it comprises the reaction of a uracil having general

formula (Ib) with an alkylating compound having general formula (VIII) according to reaction scheme 4
Scheme 4:

$$\begin{array}{c|c}
X_1 & & & \\
X_1 & & & \\
X_2 & & & \\
X_3 & & & \\
\end{array}$$
(Ib)

wherein

- X1, X2, X3 and X4 have the meanings defined above;
- R' represents a C₁-C₃ alkyl or C₁-C₃ haloalkyl group;
- L_1 represents a halogen atom, or a $R_L SO_2O-$ group wherein R_L represents a C_1-C_4 alkyl or C_1-C_4 haloalkyl group or a phenyl group possibly substituted by C_1-C_4 alkyl groups.
- 11. (previously presented): The process according to claim 10, characterized in that the reaction between the compounds having general formula (Ib) and the compound having general formula (VIII) is carried out in the presence of one or more inert organic solvents and in the presence of a base, preferably an inorganic base, at a temperature ranging from

- -10°C to the boiling point of the reaction mixture.
- 12. (withdrawn): The process according to claim 8, characterized in that the reaction is carried out in a biphasic system using water as solvent and an organic solvent immiscible with water, in the presence of phase transfer catalysts.
- 13. (withdrawn): The process for the preparation of compounds having general formula (I) according to claim 1, wherein G=O, compounds (Id), characterized in that it comprises a first reaction between a substituted aniline having formula (IV) and a chloroformiate or a carbonate having formula (IX) to give a carbamate having formula (X) and a second reaction wherein the carbamate is converted into the compounds having general formula (Id) by cyclocondensation with a 3-aminocrotonate having general formula (III), according to reaction scheme 5:

Scheme 5:

wherein

- X_1 , X_2 , X_3 , X_4 and R have the meanings defined above;
- L₅ represents a halogen atom or a OR₁₄ group;
- R_{13} and R_{14} represent a C_1-C_4 alkyl or C_1-C_4 haloalkyl group or a phenyl group possibly substituted by C_1-C_4 alkylgroups.
- 14. (withdrawn): The process according to claim 13, characterized in that the first reaction is carried out in the presence of an inert organic solvent, at a temperature ranging from -10° C to the boiling point of the mixture

itself, in the presence of an organic or inorganic base, in a quantity varying from 1 to 1.5 moles per mole of aniline (IV), with a quantity of compound having formula (IX) varying from 1 to 1.5 moles per mole of aniline (IV).

15. (withdrawn): The process according to claim 13, characterized in that the cyclo-condensation reaction of the carbamate having general formula (X) with the 3-aminocrotonate having general formula (III) is carried out in the presence of an inert organic solvent and in the presence of an organic or preferably inorganic base, at a temperature ranging from -20°C to the boiling point of the reaction mixture.

16. (withdrawn): The process according to claim 10, characterized in that the compounds having general formula (Ib) are prepared starting from an aniline having general formula (IV) by reaction with a β -ketoester having general formula (XII), to give an anilide having general formula (XIII), then converted into the intermediate of general formula (XIV) by amination with ammonia or ammonium salts, said intermediate being converted into the compounds of general formula (Ib) by cyclization with a compound of general formula (XV), such as phosgene, or diphosgene according to the reaction scheme 6

Scheme 6:

wherein:

- X_1 , X_2 , X_3 and X_4 have the meanings defined above;
- R_{13} represents a C_1 - C_4 alkyl or haloalkyl group or a phenyl group possibly substituted by C_1 - C_4 alkyl groups;
- L_6 and L_7 , having the same or different meaning, represent a chlorine atom, a CCl₃O- group, a C_1 - C_4 alkoxy group, a phenoxy group, an imidazol-1-yl group or a 1,2,4-triazol-1-yl group.
- 17. (withdrawn): The process according to claim 16,

characterised in that the reaction between the compounds having general formula (IV) and the compounds having general formula (XII) is carried out in the presence of one or more inert organic solvents, at a temperature ranging from -10°C to the boiling temperature of the reaction mixture, using an amount of compound (XII) ranging from 1 to 3 moles per mole of aniline (IV).

- 18. (withdrawn): (currently amended) The process according to claim 17, characterised in that the reaction is carried out while distilling off compound $R_{13}OH$ formed during the reaction, alone or in mixture with the solvent used.
- The process 19. (withdrawn): (currently amended) characterised in that the 16, according to claim transformation of compounds having general formula (XIII) into compounds having general formula (XIV) is carried out in the presence of one or more inert organic solvents, at a temperature ranging from -10°C to the boiling temperature of the reaction mixture, using ammonia or an ammonium salt, in an amount ranging from 1 to 20 moles per mole of compound (XIII).
- 20. (withdrawn): The process according to claim 16, characterised in that the reaction between the compounds having general formula (XIV) and the compounds having general formula (XV) is carried out in the presence of one

or more inert organic solvents, at a temperature ranging from -10°C to the boiling temperature of the reaction mixture, using an amount of compound (XV) ranging from 1 to 5 moles per mole of compound (XIV) in the presence of a suitable organic or inorganic base, in an amount ranging from 1 to 5 moles per mole of compound (XIV).

- 21. (withdrawn): Use of uracils having general formula(I) according to claims 1, as herbicides.
- 22. (withdrawn): Use according to claim 21 for the preemergence and/or post-emergence control of monocotyledonous or dicotyledonous weeds.
- 23. (withdrawn): Method for the control of weeds in cultivated areas by the application of the compounds having general formula (I) according to claims 1.
- 24. (withdrawn): (The method according to claim 23, characterized in that the amount of compound having formula (I) to be applied varies between dosages of compounds ranging from 1g to 1000g per hectare.
- 25. (currently amended): The herbicidal composition[[s]] containing, as active principle, one or more compounds

having general formula (I) according to claim 1, possibly also as a blend of isomers.

26. (currently amended): The herbicidal composition[[s]] according to claim 25, comprising other active principles which are compatible with the compounds having general formula (I), such as and are selected from the group consisting of other herbicides, fungicides, insecticides, acaricides, and fertilizers, etc.

27. (currently amended): The herbicidal composition[[s]] according to claim [[25]] 26, characterized in that the further other herbicides are selected from: acetochlor, acifluorfen, aclonifen, AKH-7088, alachlor, alloxydim, ametryn, amicarbazone, amidosulfuron, amitrole, anilofos, asulam, atrazine, azafenidin, azimsulfuron, aziprotryne, BAY MKH 6561, beflubutamid, benazolin, benfluralin, benfuresate, bensulfuron, bensulide, bentazone, benzfendizone, benzobicyclon, benzofenap, bifenox, bilanafos, bispyribac-sodium, benzthiazuron, bromacil, bromobutide, bromofenoxim, bromoxynil, butachlor, butafenacil, butamifos, butenachlor, butralin, butroxydim, butylate, cafenstrole, carbetamide, carfentrazone-ethyl, chlomethoxyfen, chloramben, chlorbromuren, chlorbufam, chlorflurenol, chloridazon, chlorimuron, chlornitroien, chlorotoluron, chloroxuron, chlorpropham, chlorsulfuron, chlorthal, chlorthiamid, cinidon ethyl, cinmethylin, cinosulfuron, clethodim, clodinafop, clomazone, clomeprop, clopyralid, cloransulam-methyl, cumyluron (JC-940), cyanazine, cycloate, cyclosulfamuron, cycloxydim, cyhalofop-butyl, 2,4-D, 2,4-DB, daimuron, dalapon, desmedipham, desmetryn, dicamba, dichlobenil, dichlorprop, dichlorprop-P, diclofop, diclosulam, diethatyl, difenoxuron, difenzoquat, diflufenican, diflufenzopyr, dimefuron, dimepiperate, dimethachlor, dimethametryn, dimethenamid, dinitramine, dinoseb, dinoseb acetate, dinoterb, diphenamid, dipropetryn, diquat, dithiopyr, 1diuron, eglinazine, endothal, EPTC, esprocarb, ethalfluralin, ethametsulfuron-methyl, ethidimuron, ethiozin (SMY 1500), ethofumesate, ethoxyfen-ethyl (HC-252), ethoxysulfuron, etobenzanid (HW 52), fenoxaprop, fenoxaprop-P, fentrazamide, fenuron, flamprop, flamprop-M, flazasulfuron, florasulam, fluazifop, fluazifop-P, fluazolate (JV 485), flucarbazone-sodium, fluchloralin, flufenacet, flufenpyr ethyl, flumetsulam, flumicloracpentyl, flumioxazin, flumipropin, fluometuron, fluoroglycofen, fluoronitrofen, flupoxam, flupropanate, flupyrsulfuron, flurenol, fluridone, flurochloridone,

fluroxypyr, flurtamone, fluthiacet-methyl, fomesafen, furyloxyfen, glufosinate, fosamine, foramsulfuron, glyphosate, halosulfuron-methyl, haloxyfop, haloxyfop-Pmethyl, hexazinone, imazamethabenz, imazamox, imazapic, imazapyr, imazaquin, imazethapyr, imazosulfuron, indanofan, iodosulfuron, ioxynil, isopropalin, isoproturon, isouron, isoxaben, isoxachlortole, isoxaflutole, isoxapyrifop, KPPlactofen, lenacil, linuron, LS830556, MCPA, MCPAmecoprop, mecoprop-P, mefenacet, thioethyl, MCPB, mesotrione, metamitron, metazachlor, mesosulfuron, methabenzthiazuron, methazole, methoprotryne, methyldymron, metobenzuron, metobromuron, metolachlor, S-metolachlor, metosulam, metoxuron, metribuzin, metsulfuron, molinate, monalide, monolinuron, naproanilide, napropamide, naptalam, NC-330, neburon, nicosulfuron, nipyraclofen, norflurazon, orbencarb, oryzalin, oxadiargyl, oxadiazon, oxasulfuron, pebulate, oxaziclomefone, oxyfluorfen, paraquat, pendimethalin, penoxsulam, pentanochlor, pentoxazone, pethoxamid, phenmedipham, picloram, picolinafen, piperophos, pretilachlor, primisulfuron, prodiamine, profluazol, proglinazine, prometon, prometryne, propachlor, propanyl, propaquizafop, propazine, propham, propisochlor, propyzamide, prosulfocarb, prosulfuron, pyraclonil, pyraflufen-ethyl, pyrazogyl (HSA-961), pyrazolynate, pyrazosulfuron, pyrazoxyfen, pyribenzoxim, pyributicarb, pyridafol, pyridate, pyriftalid, pyriminobac-methyl, pyrithiobac-sodium, quinclorac, quinmerac, quizalofop, quizalofop-P, rimsulfuron, sethoxydim, siduron, simazine, simetryn, sulcotrione, sulfentrazone, sulfometuron-methyl, sulfosulfuron, 2,3,6-TBA, TCA-sodium, tebutam, tebuthiuron, terbacil, terbumeton, terbuthyl-azine, tepraloxydim, thiazafluron, thiazopyr, terbutryn, thenylchlor, thidiazimin, thifensulfuron-methyl, thiobencarb, tiocarbazil, tioclorim, tralkoxydim, tri-allate, triasulfuron, triaziflam, tribenuron, triclopyr, trietazine, trifloxysulfuron, trifluralin, triflusulfuronmethyl, tritosulfuron, UBI-C4874, vernolate.

- 28. (currently amended): The composition[[s]] according to claim 25, characterized in that the concentration of the active substance ranges from 1 to 90%.
- 29. A uracil compound as defined in claim 1 wherein Q is 1,2,4-oxadiazolyl.
- 30. A uracil compound as defined in claim 1 wherein Q is 5-methyl-1,2,4-oxadiazolyl.